IN THE CLAIMS

- 1. (Currently Amended) A kneadable and moldable bone-replacement material which consists of a mixture of:
- A) calcium-containing ceramic particles wherein the ceramic particles comprise a calcium-phosphate ratio having a molar Ca/P relationship between 1.0 and 2.0, wherein the calcium phosphate is selected from the following group: Dicalcium-phosphate-dihydrate (CaHPO4 x 2 H2O), dicalcium-phosphate (CaHPO4), alpha-tricalcium-phosphate (alpha-Ca3(PO4)2), beta-tricalcium-phosphate (beta-Ca3(PO4)2), calcium-deficient hydro-xylapatite (Ca9(PO4)5(HPO4)OH), hydro-xylapatite (CA10(PO4)6OH)2), carbonated apatite (Ca10(PO4)3(CO3)3(OH)2), flouride-apatite (Ca10(PO4)6(F,OH)2), chloride-apatite (Ca10 (PO4)6(Cl,OH)2), whitlockite ((Ca,Mg)3(PO4)2), tetracalcium-phosphate (Ca4(PO4)2O), oxyapatite (CA10(PO4)6O), beta-calcium-pyrophosphate (beta-Ca2(P2O7), alpha-calcium-pyrophosphate, gamma-calcium-pyrophosphate, octo-calcium-phosphate (Ca8H2(PO4)6 x 5 H2O), wherein at least 50% of the ceramic particles have a pore size between 100 and 500 micrometers, wherein a bulk density of the ceramic particles is between 0.6 g/ccm and 1.0 g/ccm, wherein the jarring density of the ceramic particles is between 0.7 g/ccm and 1.1 g/ccm and wherein an average diameter of the ceramic particles is between 100 and 250 micrometers.; and
 - B) a hydrogel or a substance that can be swelled into a hydrogel, and wherein:
 - C) the ceramic particles are of fully synthetic origin;
 - D) the individual ceramic particles have at least a partially cohesive, porous structure; and
 - E) the majority of the ceramic particles have a non-spheric shape.
- 2. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles have an angular shape.
- 3. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles have a sphericity relationship S=Dmax/Dmin a largest diameter Dmax and a smallest diameter Dmin which is larger than 1.2.

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4. (Previously Presented) The bone-replacement material in accordance with claim 3, wherein the sphericity relationship S is larger than 3.

- 5. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein at least 50% of the ceramic particles have a non-spheric shape.
- 6. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein pore size of the ceramic particles is between 1 and 500 micrometers.

7.-8. (Canceled)

- 9. (Currently Amended) The bone-replacement material in accordance with claim [[8]] 1, wherein the pore size is between 340 and 450 micrometers.
- 10. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein porosity of the ceramic particles is between 60 and 90 percent.

11.-16. (Canceled)

17. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein a share of ceramic particles of non-spheric shape is at least 60%.

18.-20. (Canceled)

21. (Currently Amended) The bone-replacement material in accordance with claim [[18]] 1, wherein ceramic particles with an average diameter of 100 to 250 micrometers are used together with those having an average diameter of 250 to 500 micrometers and/or together with those having an average diameter of 0.5 to 5.6 mm.

22.-25 (Canceled)

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26. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles consist of a mixture of different calcium-phosphates.

27. (Canceled)

- 28. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles consist of a calcium-carbonate.
- 29. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles are selected from the following group: alpha-calcium-sulfate-hemihydrate, beta-calcium-sulfate-hemihydrate, calcium-sulfate-dihydrate.
- 30. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the ceramic particles consist of a mixture of different calcium-phosphates, calcium-sulfates and/or calcium-carbonates.
- 31. (Previously Presented) The bone-replacement material in accordance with claim 1, further comprising metallic or semi-metallic ion shares as additives.
- 32. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel consists of fully synthetic substances.
- 33. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel consists of natural biological substances, preferably of plant origin.
- 34. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel consists of a biotechnologically generated substance.

RESPONSE TO RESTRICTION REQUIREMENT

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- 35. (Previously Presented) The bone-replacement material in accordance with one claim 32, wherein the hydrogel or the substance which can be swelled into a hydrogel consists of a mixture of fully synthetic, natural biological or biotechnologically generated substances.
- 36. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel contains one of the following components: a) polyamino-acids or their derivatives, preferably polylysin or gelatin; b) polysaccharides and their derivatives, preferably glycosaminoglycane or alginate; c) polylipides, fatty acids and their derivatives; d) nucleotides and their derivatives; or a combination of the components as listed in a) through d).
- 37. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel contains one of the following components: a) polymethylenoxide or its derivatives; b) polyethylene, polypropylenoxide or their derivatives; d) polyacrylate or its derivatives; or a combination of the components as listed in a) through d).
- 38. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel or the substance which can be swelled into a hydrogel consists of either a glycosaminoglycane or a proteoglycane or a mixture of those two substances.
- 39. (Previously Presented) The bone-replacement material in accordance with claim 38, wherein the glycosaminoglycane is a hyaluron-acid, chondroitinsulfate, dermatansulfate, heparansulfate, heparansulfate.
- 40. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein a concentration of the ready-to-use, hydrated hydrogel or a ready-to-use, hydrated substance which can be swellen into a hydrogel is between 0.1% and 20.0%.

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41. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein a molecular weight of the hydrogel or the substance which can be swelled into a hydrogel is larger than 300,000 Dalton and preferably larger than 500,000 Dalton.

- 42. (Previously Presented) The bone-replacement material in accordance with claim 41, wherein the molecular weight of the hydrogel or the substance which can be swelled into a hydrogel is larger than 1,000,000 Dalton and preferably larger than 1,500,000 Dalton.
- 43. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein the hydrogel is a liquid solution of a hyaluronate.
- 44. (Previously Presented) The bone-replacement material in accordance with claim 43, wherein the liquid solution of the hydrogel contains less than 99% water.
- 45. (Previously Presented) The bone-replacement material in accordance with claim 43, wherein the liquid solution of the hydrogel contains less that 96.5% water.
- 46. (Previously Presented) The bone-replacement material in accordance with claim 43, wherein the molecular weight of the hyaluron-acid used is larger than 1.5×10^6 Dalton.
- 47. (Previously Presented) The bone-replacement material in accordance with claim 43, wherein the molecular weight of the hyaluron-acid used is between 0.5×10^6 and 1.0×10^6 Dalton.
- 48. (Previously Presented) The bone-replacement material in accordance with claim 43, wherein the molecular weight of the hyaluron-acid used is smaller than 1×10^6 and preferably smaller than 0.5×10^6 Dalton.

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- 49. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein a specific gravity of the calcium-containing, porous ceramic particles is between 0.5 and 1.0 g/ccm.
- 50. (Previously Presented) The bone-replacement material in accordance with claim 1, wherein a weight relationship A/B between the hydrated hydrogel and the calcium-containing ceramic particles is larger than 0.2.
- 51. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is between 0.2 and 0.5.
- 52. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is between 0.5 and 0.9.
- 53. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is between 0.9 and 1.3.
- 54. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is between 1.3 and 2.0.
- 55. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is between 2 and 5.
- 56. (Previously Presented) The bone-replacement material in accordance with claim 50, wherein the weight relationship A/B is larger than. 5.